Billing Code 4410-09-M

DEPARTMENT OF JUSTICE

DRUG ENFORCEMENT ADMINISTRATION

MANUFACTURER OF CONTROLLED SUBSTANCES

NOTICE OF APPLICATION

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on December 15, 2011, Pharmagra Labs, Inc., 158 McLean Road, Brevard, North Carolina 28712, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Pentobarbital (2270), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed substance for analytical research and clinical trials.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR § 1301.33(a).

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register

Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than [insert date 60 days from date of publication].

Joseph T. Rannazzisi
Deputy Assistant Administrator
Office of Diversion Control
Drug Enforcement Administration

DATED: January 30, 2012

[FR Doc. 2012-2608 Filed 02/03/2012 at 8:45 am; Publication Date: 02/06/2012]